effectiveness, including what will be evaluated and analyzed, who will perform the evaluation and the time-frame.

6. Collaboration (15 points)

The extent to which the applicant documents evidence of collaboration and experience with partners.

7. Budget (Not scored)

The extent to which the budget is clearly explained, adequately justified, reasonable, sufficient for the proposed project activities, and consistent with the intended use of the cooperative agreement funds.

H. Other Requirements

Technical Reporting Requirements

Provide CDC with original plus two copies of:

1. An annual progress report.
2. A financial status report, no more than 90 days after the end of the budget period; and
3. A final financial status and performance report, no more than 90 days after the end of the project period.

The following additional requirements are applicable to the program. For a complete description of each see Addendum 1 in the application package.

AR–5 HIV Program Review Panel Requirements
AR–10 Smoke-Free Workplace Requirements
AR–11 Healthy People 2000
AR–12 Lobbying Restrictions
AR–13 Prohibition on Use of CDC Funds for Certain Gun Control Activities
AR–14 Accounting System Requirements
AR–15 Proof of Non-profit Status

I. Authority and Catalog of Federal Domestic Assistance Number

This program is authorized under sections 301(a) and 317(k)2) of the Public Health Service Act (42 U.S.C. 241(a) and 247b(k)2)), as amended.

J. Where To Obtain Additional Information

This and all other CDC Announcements may be found and downloaded from the CDC homepage. Internet address: http://www.cdc.gov (click on funding). To receive additional written information and to request an application kit, call 1–888–GRANTS4 (1–888–472–6874). You will be asked to leave your name and address and will be instructed to identify the Announcement number of interest, 99153.

If you have questions after reviewing the contents of all the documents, business management technical assistance may be obtained from: Joanne Wojcik, Grants Management Specialist, Grants Management Branch, Procurement and Grants Office, Announcement 99153, Centers for Disease Control and Prevention, 2920 Brandywine Road, Suite 3000, Mailstop E–13, Atlanta, GA 30341–4146; Telephone: (770) 488–2717; E-mail address: jcw6@cdc.gov.

For program technical assistance contact: Nancy Chalmers, M.P.A., Centers for Disease Control and Prevention, Office of Program Planning and Evaluation, 1600 Clifton Road, Mailstop D–24, Atlanta, GA 30333; Telephone: (404) 639–7085; E-mail address: npc1@cdc.gov.

Dated: June 10, 1999.

John L. Williams,

Director, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 99–15218 Filed 6–15–99; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Ophthalmic Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Ophthalmic Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA's regulatory issues.

Date and Time: The meeting will be held on July 22 and 23, 1999, 8 a.m. to 5:30 p.m.

Location: Hilton Hotel, Salons A and B, 620 Perry Pkwy., Gaithersburg, MD.

Contact Person: Sara M. Thornton, Center for Devices and Radiological Health (HFZ–460), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12396. Please call the Information Line for up-to-date information on this meeting.

Agenda: On July 22, 1999, the committee will discuss, make recommendations, and vote on a premarket approval application (PMA) for the correction of myopia with and without astigmatism using laser in-situ keratomileusis (LASIK). The committee will also discuss, make recommendations, and vote on a holmium laser for the correction of hyperopia using laser thermal keratomileusis. On July 23, 1999, the committee will discuss, make recommendations, and vote on a soft acrylic intraocular lens for the visual correction of aphakia after cataract extraction. The committee will also discuss, make recommendations, and vote on a PMA for the correction of myopia with and without astigmatism using LASIK.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by July 12, 1999. Oral presentations from the public will be scheduled between approximately 8:30 a.m. and 9 a.m. on July 22 and 23, 1999. Near the end of the committee deliberations on each PMA, a 30-minute open public session will be conducted for interested persons to address issues specific to the submission before the committee. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before July 12, 1999, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: June 9, 1999.

Michael A. Friedman,

Deputy Commissioner for Operations.

[FR Doc. 99–15183 Filed 6–15–99; 8:45 am]

BILLING CODE 4160–01–F